depleted of activity of proteins that bind polyadenylate;

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ii) a source of ATP; and

iii) [said] an exogenous target RNA sequence.

In claim 2, lines 4, 2 and 3, please change the occurrences of "turnover" to -- deadenylation and degradation --.

Please cancel claims 7 and 8.

In claim 9, line 1, please change "8" to - - 1 - -.

In claim 21, lines 5 and 6, please change the occurrences of "turnover" to

- - deadenylation and degradation - -.

In claim 27, line 1, please change "turnover" to - - deadenylation and degradation - -.

In claim 33, lines 7 and 8, please change both occurrences of "turnover" to - - deadenylation and degradation - -.

Please rewrite claim 38 as follows.

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(Amended) The method of claim 36 [33] wherein said determining the extent of deadenylation and degradation [turnover] of said target RNA sequence comprises determining the extent of degradation of said labeled target RNA.

Please amend claim 46 as follows.

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(Amended) A method for identifying an agent capable of modulating <u>regulated</u> [the] deadenylation of a target RNA sequence comprising

- (A) providing a [the] system that recapitulates regulated RNA deadenylation of an exogenously-added preselected target RNA sequence comprising
 - i) a cytoplasmic extract supernatant from a 100,000 x g, 1 hour centrifugation isolated from eukaryotic cells or tissues, said extract depleted of activity of proteins that bind polyadenylate;

- ii) said target RNA sequence [the system of claim 1 in the absence of a nucleotide triphosphate];
- (B) introducing said agent into said system;
- (C) monitoring the deadenylation of said target RNA sequence in said system; and

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(D) identifying an agent able to modulate the extent of said deadenylation as capable of modulating the <u>regulated</u> deadenylation of said target RNA sequence.

In claim 47, line 1, please change "the" to - - regulated - -.

In claim 51, line 6, please change "turnover" to - - deadenylation and degradation - -.

Please amend claim 53 as follows:

Twice Amended) A kit for monitoring the stability of a preselected exogenous target

RNA sequence under conditions capable of recapitulating regulated RNA deadenylation

and degradation [turnover], said kit comprising:

- (a) <u>a cytoplasmic</u> [cell] extract <u>supernatant from a 100,000 x g, 1 hour</u> <u>centrifugation, said extract</u> depleted of activity of proteins that bind polyadenylate;
- (b) other reagents; and
- (c) directions for use of said kit.

Remarks

Claims 1, 2 and 4-55 are pending in the application. By way of the foregoing amendment, the subject matter of claims 7 and 8 have been incorporated into claims 1 and 46, the latter which has been rewritten as an independent claim. As a result, claims 1, 2, 4-6 and 9-55 are pending in